FDA

MAUDE Adverse Event Report: EPIC MYCHCART CPOE/EHR

EPIC MYCHCART CPOE/EHR

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Event Date 06/01/2015
Event Type Injury
Event Description

The drop down menu to specify route of administration of medications is a source of innumerable errors. The first option is im and is frequently selected because of its location and close resemblance to iv. It is a flaw that has been carried on by the vendor for years, endangering hundreds of patients, if not thousands over time.

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Brand Name MYCHCART
Type of Device CPOE/EHR
Manufacturer *(Section D)* EPIC
Verona WI 53593
MDR Report Key 4834779
Report Number MW5043001
Device Sequence Number 1
Product Code LNX[24]
Report Source Voluntary
Reporter Occupation Physician
Type of Report Initial
Report Date 06/03/2015
***1* Device Was Involved in the Event**
***1* Patient Was Involved in the Event**
Date FDA Received 06/03/2015
Is This An Adverse Event Report? No
Is This A Product Problem Report? Yes
Device Operator Health Professional
Is The Reporter A Health Professional? Yes
Is this a Reprocessed and Reused Single-Use Device? No

Patient TREATMENT DATA
Date Received: 06/03/2015 Patient Sequence Number: 1

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Page Last Updated: 01/31/2019

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